FDA 510(k) Premarket Notification

K872838 NORFOLK NORPORT (TM) -SP (SKIN PARALLEL)

CLEARED

Aug 25, 1987
36 days to decision

K872838 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k872838/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jul 20, 1987
Decision date	Aug 25, 1987
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Norfolk Medical Products, Inc.**
Location	Walker, MI, US
Contact	MICHAEL J DALTON
510(k) history	20 submissions · 20 cleared · 1983-2020